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2. The method of claim 1 which increases cerebral blood flow and reduces infarct size in the mammal.

4. The method of claim 3 wherein the anti-CD18 antibody fragment is a F(ab')<sub>2</sub>.

5. The method of claim 1 wherein the anti-CD18 antibody is humanized.

6. The method of claim 1 wherein the anti-CD18 antibody is administered to the mammal by bolus dosage.

7. The method of claim 1 wherein the anti-CD18 antibody is administered intravenously.

8. The method of claim 1 wherein the anti-CD18 antibody is administered via continuous infusion.

9. The method of claim 1 wherein the anti-CD18 antibody is administered to the mammal at a time between about 15 minutes to about 20 hours from the onset of focal ischemic stroke.

10. The method of claim 9 wherein the anti-CD18 antibody is administered to the mammal at a time between about 45 <sup>minutes</sup> mins to about 5 hours from the onset of focal ischemic stroke.

11. The method of claim 1 wherein the anti-CD18 antibody is humanized H52 antibody.

12. The method of claim 11 wherein the H52 antibody is a F(ab')<sub>2</sub>.

13. The method of claim 3 wherein the anti-CD18 antibody fragment is fused to a salvage receptor binding epitope.

14. The method of claim 1 wherein the mammal is a human.

15. An article of manufacture, comprising:

a container;

a label on said container; and

a composition comprising an active agent contained within said container; wherein the composition is effective for increasing cerebral blood flow or reducing infarct size in focal ischemic stroke caused by obstruction of a main cerebral artery, the label on said container indicates that the composition can be used for treating stroke and the active agent in said composition is an antagonist anti-CD18 antibody.

16. The article of manufacture of claim 15 further comprising instructions for administering the anti-CD18 antibody to a mammal to increase cerebral blood flow or reduce infarct size in focal ischemic stroke.

17. A kit, comprising:

a first container, a label on said container, and a composition comprising an active agent contained within said container; wherein the composition is effective for increasing cerebral blood flow or reducing infarct size in focal ischemic stroke caused by obstruction of a main cerebral artery, the label on said container indicates that the composition can be used for treating stroke, and the active agent in said composition is an antagonist anti-CD18 antibody;

a second container comprising a pharmaceutically-acceptable buffer; and  
instructions for using the anti-CD18 antibody to increase cerebral blood flow or reduce infarct size in focal ischemic stroke.